Attorney Docket No: 23540-10616/US Client Ref: UC-2001-F10-2

USSN: 09/927,315

AMENDMENTS TO THE CLAIMS (NONE)

This listing of claims will replace all prior versions and listings of claims in the application:

- 1-48. (Canceled)
- 49. (CURRENTLY AMENDED) A method of identifying a compound that activates or inhibits sweet taste signal transduction in taste cells, the method comprising the steps of:
- (i) contacting the compound with a cell expressing a sweet taste receptor comprising a T1R3 polypeptide and a T1R2 polypeptide, wherein the T1R3 polypeptide has a greater than 90% amino acid sequence identity to SEQ ID NO: 20, 23, or 25 greater than 93% amino acid sequence identity to SEQ ID NO: 20 or greater than 93% amino acid sequence identity to SEQ ID NO: 23; and wherein the T1R2 polypeptide has a greater than 90% 92% amino acid sequence identity to SEQ ID NO: 7 or 8 SEQ ID NO:8; and
- (ii) determining the functional effect of the compound upon the receptor, wherein said functional effect is binding to the receptor or having an effect on the activity of the receptor, thereby identifying a compound that activates or inhibits sweet signal transduction,

wherein the sweet taste receptor specifically binds a sweet compound, and at least one of the T1R3 and T1R2 polypeptides is recombinant.

- 50. (Previously presented) The method of claim 49, wherein the T1R2 polypeptide and the T1R3 polypeptide are non-covalently linked.
- 51. (Previously presented) The method of claim 49, wherein the T1R2 polypeptide and the T1R3 polypeptide are covalently linked.
 - 52-55. (Canceled)
- 56. (**CURRENTLY AMENDED**) The method of claim 49, wherein the T1R3 polypeptide has an amino acid sequence of SEQ ID NO: 20, 23, or 25 <u>SEQ ID NO: 20 or 23</u>.
- 57. (**CURRENTLY AMENDED**) The method of claim 49, wherein the T1R2 polypeptide has an amino acid sequence of SEQ ID NO: 7 or 8 SEQ ID NO: 8.

Attorney Docket No: 23540-10616/US Client Ref: UC-2001-F10-2

USSN: 09/927,315

58. (Previously presented) The method of claim 49, wherein the T1R3 and T1R2

polypeptides are both recombinant.

59-66. (Canceled)

67. (Previously presented) The method of claim 49, wherein the functional effect is

determined by measuring ligand binding to the receptor.

68. (Canceled)

69. (Previously presented) The method of claim 49, wherein the functional effect is a

chemical or phenotypic effect.

70. (Previously presented) The method of claim 49, wherein the functional effect is

determined by measuring changes in intracellular cAMP, IP3, or Ca²⁺.

71. (Previously presented) The method of claim 49, wherein the cell is a mammalian

cell.

72. (Previously presented) The method of claim 71, wherein the cell is a human cell.

73-74. (Canceled)

75. (CURRENTLY AMENDED) A method of identifying a compound that

activates or inhibits sweet taste signal transduction in taste cells, the method comprising the steps

of

(i) contacting the compound with a cell expressing a sweet taste receptor comprising a

T1R3 polypeptide and a T1R2 polypeptide, wherein the T1R3 polypeptide has a greater than

90% amino acid sequence identity 93% amino acid sequence identity to SEQ ID NO:20; and

wherein the T1R2 polypeptide has a greater than 90% amino acid sequence identity 92% amino

acid sequence identity to SEQ ID NO:8; and

(ii) determining the functional effect of the compound upon the receptor, wherein said

functional effect is binding to the receptor or having an effect on the activity of the receptor,

thereby identifying a compound that activates or inhibits sweet signal transduction,

wherein the sweet taste receptor specifically binds a sweet compound, and at least one of

the T1R3 and T1R2 polypeptides is recombinant.

Attorney Docket No: 23540-10616/US

Client Ref: UC-2001-F10-2

USSN: 09/927,315

76. (Previously presented) The method of claim 75, wherein the T1R2 polypeptide has the amino acid sequence of SEQ ID NO:8.

77-78. (Canceled)